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⑯ **Laminated structure.**

⑯ A laminated structure comprising, laminated in the order
stated,

- (A) a layer of a polyester,
- (B) a layer of a modified ethylene/alpha-olefin random
copolymer, the modified copolymer containing about 0.01 to
10% by weight, based on the trunk ethylene/alpha-olefin ran-
dom copolymer, of an unsaturated carboxylic acid or its deriva-
tive grafted thereto, and having an ethylene content of 75 to
95 mole%, a melt flow rate of 0.1 to 50 g/10 min., a density
of 0.850 to 0.905 g/cm³ and a crystallinity of less than 30%,
and
- (C) a layer of a saponified olefin/vinyl acetate copolymer
or a layer of a polyamide resin.

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LAMINATED STRUCTURE

This invention relates to a laminated structure composed of a plurality of polymer layers. More specifically, it relates to a laminated structure which comprises a polyester layer and a saponified olefin/vinyl acetate copolymer layer or a polyamide layer and has transparency and resistance to permeation to gases and excellent interfacial adhesion, particularly excellent interfacial adhesion after stretching.

Polyester resins typified by polyethylene terephthalate (to be sometimes referred to as PET) are extensively used as films, sheets and packaging materials such as containers by utilizing their excellent properties such as high mechanical strength, rigidity, thermal resistance, chemical resistance, oil resistance and transparency. However, the polyester resins are not without defects. For example, they have the defect that containers made of the polyester resins cannot be sufficiently heat-set, and they have no sufficient resistance to gas permeation. In particular, the insufficient gas permeation resistance is attributed to the inherent properties of the polyester resins, and is difficult to overcome by molding techniques.

It may also be possible to improve the gas permeation resistance of the polyester resins by laminating a resin having better gas permeation resistance than the polyester resins, such as polyvinylidene chloride, a saponification product of olefin/vinyl acetate copolymer, or polyamides. Such resins, however, have poor adhesion to the polyester resins. Even a three-layer structure composed of two layers of a polyester resin and a layer of such a resin having high gas permeation resistance interposed between them is likely to undergo partial delamination during stretching or during use and decrease in gas permeation resistance.

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A composition composed of 60 to 95% by weight of a polyolefin modified with an unsaturated carboxylic acid or its anhydride and 40 to 5% by weight of a rubbery material was proposed as a composition having excellent adhesion to a polyester resin, a saponified ethylene/vinyl acetate copolymer, or a polyamide resin (U. S. Patent No. 4,058,647). This composition has an excellent initial adhesion strength, but has reduced transparency. If a laminated structure of the polyester resin and this composition is stretched in order to improve the rigidity of the polyester resin, the adhesion strength between the two layers is greatly reduced, and it has been found that the resulting laminate gives rise to problems when used as a packaging.

It is an object of this invention to provide a novel laminated structure.

Another object of this invention is to provide a novel laminated structure having a polyester layer and a saponified olefin/vinyl acetate copolymer layer or a polyamide layer and an interlayer having excellent adhesion strnegrth to both of these layers.

Still another object of this invention is to provide a novel laminated structure having a polyester layer and a saponified olefin/vinyl acetate copolymer layer or a polyamide layer and an interlayer which does not impair the transparency of these two layers.

Yet another object of this invention is to provide a novel laminated structure having a polyester layer and a saponified olefin/vinyl acetate copolymer layer or a polyamide layer and an interlayer which even after stretching, retains excellent adhesión to the two layers.

A further object of this invention is to provide a laminated structure having a polyester layer and a saponified olefin/vinyl acetate copolymer layer or a polyamide layer which are bonded through an interlayer of

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a modified ethylene/alpha-olefin random copolymer which is obtained by grafting an unsaturated carboxylic acid or its derivative and has a very low crystallinity or is amorphous.

5. Additional objects of the invention along with its advantages will become apparent from the following description.

According to this invention, the above objects and advantages of the invention are achieved by a laminated 10 structure comprising, laminated in the order stated,

- (A) a layer of polyester,
- (B) a layer of a modified ethylene/alpha-olefin random copolymer, the modified copolymer containing about 0.01 to 10% by weight, based on the trunk ethylene/alpha-15 olefin random copolymer, of an unsaturated carboxylic acid or its derivative grafted thereto, and having an ethylene content of 75 to 95 mole%, a melt flow rate of 0.1 to 50 g/10 min., a density of 0.850 to 0.905 g/cm³ and a crystallinity of less than 30%, and
- 20 (C) a layer of a saponified olefin/vinyl acetate copolymer or a polyamide resin.

The thermoplastic polyester (A) denotes polyesters composed of units of dihydroxy compounds selected from aliphatic glycols such as ethylene glycol, propylene 25 glycol, 1,4-butanediol, neopentyl glycol and hexamethylene glycol, alicyclic glycols such as cyclohexanediethanol, aromatic dihydroxy compounds such as bisphenol or mixtures of at least two types of these and units of dicarboxylic acids selected from aromatic dicarboxylic acids such as 30 terephthalic acid, isophthalic acid and 2,6-naphthalenedi-carboxylic acid, aliphatic dicarboxylic acids such as oxalic acid, succinic acid, adipic acid, sebamic acid and undecanedicarboxylic acid, alicyclic dicarboxylic acids such as hexahydroterephthalic acid, or mixtures of at 35 least two types of these. These polyesters may be modified with small amounts of trihydric or higher polyhydroxy

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compounds such as triols or tricarboxylic acids, or polycarboxylic acids so long as the polyesters remain thermoplastic. Specific examples of the thermoplastic polyesters are polyethylene terephthalate, polybutylene 5 terephthalate, and polyethylene isophthalate terephthalate copolymer.

The saponified olefin/vinyl acetate copolymer constituting the layer (C) of the laminated structure of the invention is a saponification product of an olefin/vinyl acetate copolymer having an olefin content of 15 to 10 60 mole%, preferably 25 to 50 mole%. The saponification product has a degree of saponification of at least 50%, more preferably at least 90%. If the olefin content is less than 15 mole%, the saponified copolymer tends to be 15 not entirely satisfactory in respect of resistance to thermal decomposition, melt-processability, stretch-ability and water resistance (moisture absorption and 20 swellability). If the olefin content exceeds 60 mole% or the degree of saponification is less than 50%, the copolymer has poor resistance to gas permeation.

Alpha-olefins having 1 to 18 carbon atoms are preferred as the olefin for composing the saponified olefin/vinyl acetate copolymer. Examples include ethylene, propylene, 1-butene, 1-hexene, 4-methyl-1- 25 pentene, 1-octene, 1-decene, 1-tetradecene and 1-octadecene.

Saponified olefin/vinyl acetate copolymer is especially preferred as the saponified olefin/vinyl acetate copolymer in view of its mechanical strength and 30 processability.

Examples of the polyamide resin-(C) constituting the laminated structure of the invention include polyamides obtained by polycondensation of aliphatic, alicyclic or aromatic diamines such as hexamethylenediamine, decamethylenediamine, dodecamethylenediamine, 35 2,2,4- or 2,4,4-trimethylhexamethylenediamine, 1,3- or

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1,4-bis(aminomethyl)cyclohexane, bis(p-aminocyclohexyl-methane) and m- or p-xylylenediamine and aliphatic, alicyclic or aromatic dicarboxylic acids such as adipic acid, suberic acid, sebacic acid, cyclohexanedicarboxylic acid, terephthalic acid and isophthalic acid; polyamides obtained by polycondensation of aminocarboxylic acids such as epsilon-aminocaproic acid and 11-aminoundecanoic acid; polyamides obtained by ring-opening polymerization of lactams such as epsilon-caprolactam and omega-lauro-lactam; copolyamides composed of two or more of these components; and mixtures of these polyamides. Specific examples are nylon 6, nylon 66, nylon 610, nylon 9, nylon 11, nylon 12, nylon 6/66, nylon 66/610 and nylon 6/11. There is no restriction on the molecular weights of these polyamides. Usually polyamides having a relative viscosity η_r (measured in 98% sulfuric acid in accordance with JIS K-6819) of at least 0.5, preferably at least 2.0, are conveniently used in this invention.

The laminated structure of this invention comprises the layer (B) of a modified ethylene/alpha-olefin random copolymer as an interlayer for firmly bonding the layers (A) and (C).

The modified ethylene/alpha-olefin random copolymer (to be sometimes referred to simply as the modified copolymer) contains about 0.01 to 10% by weight, preferably 0.1 to 5% by weight, based on the ethylene/alpha-olefin random copolymer as a trunk, of an unsaturated carboxylic acid or its derivative. If the amount of the unsaturated carboxylic acid or its derivative grafted is less than 0.01% by weight, the adhesion of the interlayer to the polyester layer (A) and the saponified olefin/vinyl acetate copolymer layer or the polyamide layer (C) cannot be improved. On the other hand, if it exceeds 10% by weight, the modified copolymer is partly crosslinked and is inferior in processability, transparency and adhesion strength.

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The modified copolymer has an ethylene content of 75 to 95 mole%, preferably 77 to 93 mole%.

The modified copolymer also has a melt flow rate (MFR_2 in accordance with ASTM D1238L) of 0.1 to 50 g/10 min., preferably 0.2 to 20 g/10 min.

The modified copolymer has a density of 0.850 to 0.905 g/cm³, preferably 0.860 to 0.900 g/cm³.

Finally, the modified copolymer has a crystallinity of less than 30%, preferably less than 10%, more 10 preferably less than 7%. If the MFR_2 is outside the above-specified range, the viscosity of the modified copolymer is too high or too low and has poor processability and adhesion strength.

If the modified copolymer has a density of more 15 than 0.905 g/cm³ and a crystallinity, determined by X-ray, of more than 30%, the modified copolymer is inferior in adhesion strength after it has been subjected to stretching.

Alpha-olefins having 3 to 20 carbon atoms are 20 suitable as the alpha-olefin constituting the modified copolymer. Specific examples are propylene, 1-butene, 1-hexene, 4-methyl-1-pentene, 1-octene, 1-decene, 1-tetradecene and 1-octadecene. These alpha-olefins may be used singly or in combination.

25 The modified copolymer of layer (B) can be produced by grafting the unsaturated carboxylic acid or its derivative to an ethylene/alpha-olefin random copolymer. The ethylene/alpha-olefin random copolymer before modification usually has a melt flow rate (MFR_2) of 0.1 30 to 70 g/10 min., and has the same density, ethylene content and crystallinity as the modified copolymer.

Examples of the unsaturated carboxylic acid or its derivative to be grafted include unsaturated carboxylic acids such as acrylic acid, maleic acid, fumaric acid, 35 tetrahydrophthalic acid, itaconic acid, citraconic acid, crotonic acid, isocrotonic acid, and Nadic acid[®] (endo-

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cis-bicyclo[2.2.1]hept-5-ene-2,3-dicarboxylic acid); and derivatives thereof such as acid halides, amides, imides, anhydrides and esters. Specific examples of the derivatives are malenyl chloride, maleimide, maleic anhydride, 5 citraconic anhydride, monomethyl maleate, dimethyl maleate, and glycidyl maleate. Among these, the unsaturated dicarboxylic acids and their anhydrides are preferred. Maleic acid, Nadic acid[®] and their anhydrides are especially preferred.

10 Various known methods can be used to produce the modified copolymer by grafting a grafting monomer selected from the unsaturated carboxylic acids and their derivatives to the unmodified ethylene/alpha-olefin random copolymer. For example, the unmodified ethylene/ 15 alpha-olefin random copolymer is melted, and the grafting monomer is added to it and graft-copolymerized. Alternatively, the unmodified ethylene/alpha-olefin random copolymer is dissolved in a solvent, and the grafting monomer is added and graft-copolymerized. In any case, 20 the reaction is preferably carried out in the presence of a radical initiator in order to perform graft-copolymerization with good efficiency. The grafting reaction is carried out usually at a temperature of 60 to 350°C. The proportion of the radical initiator is usually 0.001 25 to 1 part by weight per 100 parts by weight of the unmodified ethylene/alpha-olefin random copolymer. The radical initiator may be an organic peroxide, an organic perester or an azo compound. Specific examples include benzoyl peroxide, dichlorobenzoyl peroxide, dicumyl peroxide, 30 di-tert-butyl peroxide, 2,5-dimethyl-2,5-di(peroxybenzoate)hexyne-3, 1,4-bis(tert-butylperoxyisopropyl)benzene, lauroyl peroxide, tert-butyl peracetate, 2,5-dimethyl-2,5-di(tert-butylperoxy)hexyne-3, 2,5-dimethyl-2,5-di-(tert-butylperoxy)hexane, tert-butyl perbenzoate, tert- 35 butyl perphenylacetate, tert-butyl perisobutyrate, tert-butyl per-sec-octoate, tert-butyl perpivalate, cumyl

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perpivalate, tert-butyl perdiethylacetate, azobisisobutyronitrile and dimethyl azoisobutyrate. Among these, dialkyl peroxides such as dicumyl peroxide, di-tert-butyl peroxide, 2,5-dimethyl-2,5-dimethyl-2,5-di(tert-butylperoxy)hexane and 1,4-bis(tert-butylperoxyisopropyl)benzene are preferred.

The modified copolymer used in this invention may be mixed with an unmodified ethylene/alpha-olefin random copolymer to form the modified copolymer layer (B) of the laminated structure of the invention so long as the amount of the unsaturated carboxylic acid or its derivative grafted in the mixture is in the range of 0.01 to 10% by weight. Furthermore, the modified copolymer used in this invention may be mixed with a modified or unmodified ethylene/alpha-olefin random copolymer having a crystallinity of more than 30% to form the modified copolymer layer (B) of the laminated structure of the invention so long as the crystallinity of the mixture is less than 30%.

20 A heat stabilizer, a weatherability stabilizer, an antistatic agent, a pigment, a dye, a corrosion inhibitor, etc., may be incorporated in one or more of the layers (A), (B) and (C) forming the laminated structure of this invention so long as the additives do not impair 25 the objects of this invention.

The laminated structure of this invention may, for example, be in the form of a film, sheet or bottle.

The laminated structure of this invention may be produced, for example, by a method which comprises 30 melting the polymers for the three layers in different extruders, feeding the molten polymers into a die of a three-layer structure, and co-extruding them with the modified copolymer layer (B) as an interlayer, or a method (so-called sandwich laminate method) which comprises forming the polyester layer (A) and the saponified 35 olefin/vinyl acetate copolymer layer or the polyamide

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resin layer (C) separately, and melt-extruding the modified copolymer as an interlayer into between the layers (A) and (C). If the laminated structure to be produced is in the form of a bottle, a keg, a pipe or a tube, the 5 co-extrusion molding method can be used advantageously. Furthermore, the use of the co-extrusion molding method is preferred because it can easily give a high interfacial adhesion.

The laminated structure of this invention may 10 be stretched monoaxially or biaxially in order to improve rigidity, transparency, mechanical strength, etc. or impart shrinkability. There is no particular restriction on the stretching conditions. Generally, the stretching is carried out at a temperature of 70 to 150°C at a 15 stretch ratio of 2 to 10 times in the case of monoaxial stretching and at a stretch ratio of 2 to 10 times in the longitudinal direction and 2 to 10 times in the transverse direction in the case of biaxial stretching.

The thickness of each of the layers of the 20 laminated structure can be properly determined depending upon the end use of the laminated structure. Usually, if the laminated structure is a film or sheet, the polyester resin layer (A) preferably has a thickness of 0.02 to 5 mm, and the layers (B) and (C) preferably have a thickness of 0.01 to 1 mm. If the laminated structure is a 25 container, the polyester resin layer (A) preferably has a thickness of 0.1 to 5 mm, and the layer (B) or (C) preferably has a thickness of 0.01 to 1 mm.

The laminated structure of this invention may 30 comprise three layers (A), (B) and (C), or may comprise 5 or more layers additionally having another kinds of polymer layers such as polyolefin. An example of the multilayer laminate is one comprising, for example, the polyester layer (A)/the modified random copolymer layer 35 (B)/the saponified olefin-vinyl acetate copolymer layer or the polyamide resin layer (C)/the modified random

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copolymer layer (B)/a polypropylene layer.

Depending upon its use, the laminated structure of this invention may be a 5-layer structure comprising three types of polymer, which, for example, comprises the 5 polyester resin layer (A)/modified random copolymer layer (B)/saponified olefin-vinyl acetate copolymer layer or the polyamide resin layer (C)/modified random copolymer layer (B)/polyester resin layer (A).

The laminated structure of this invention can 10 be used, for example as packaging film or bottles for foods and medicines since it has transparency, flavor retaining property and heat resistance which are the characteristic features of the polyester and resistance to gas permeation which is the characteristic feature of 15 the saponified olefin/vinyl acetate copolymer or the polyamide resin, and possesses a high interfacial adhesion in the initial stage and after stretching.

The following examples illustrate the present invention in greater detail. It should be understood 20 that unless the invention departs from its scope described and claimed herein, it is not to be limited in any way to these examples.

EXAMPLE 1

A three-layer case sheet was formed under the 25 following conditions by using maleic anhydride-grafted ethylene/propylene random copolymer (MAH-EPR-1; MFR₂ 0.5 g/10 min., the amount of maleic anhydride grafted 0.5 g/100 g polymer, ethylene content 80 mole%, density 0.865 g/cm³, crystallinity 4%), polyethylene terephthalate 30 (PET; tradename Mitsui PET J 135, a product of Mitsui PET Resin Co., Ltd.), and a saponified product of ethylene/vinyl acetate copolymer (EVOH; MFR₂ 1.3 g/10 min., density 1.19 g/cm³, ethylene content 32 mole%; tradename Kuraray Eval EP-F, a product of Kuraray Inc.).

35 Film layer construction:

PET/MAH-EPR-1/EVOH=120/60/120 micrometers

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Extruder:

- 40 mmØ extruder 280°C (for PET)
- 40 mmØ extruder 210°C (for MAH-EPR-1)
- 40 mmØ extruder 210°C (for EVOH)

5 Molding speed: 10 meters/min.

The interfacial adhesion strength (F_1 , g/15 mm width) between the PET layer and the MAH-EPR-1 layer of the resulting three-layer sheet, and the interfacial adhesion strength (F_2 , g/15 mm width) between the EVOH 10 layer and the MAH-EPR-1 layer were measured by T-peel method at a peeling speed of 300 mm/min. The transparency of the resulting sheet was evaluated by using a haze meter. The three layer sheet was heated at 80°C for 10 minutes, and at this temperature stretched simultaneously 15 to three times longitudinally and to three times transversely to prepare a three-layer stretched sheet. Subsequently, the sheet was cooled to room temperature, and the interfacial adhesion strength (F_3 , g/15 mm width) between the PET layer and the MAH-EPR-1 layer and the 20 interfacial adhesion strength (F_4 , g/15 mm width) between the EVOH layer and the MAH-EPR-1 layer in the resulting stretched sheet were measured by T-peel at a peeling speed of 300 mm/min. The transparency of the three-layer stretched sheet was evaluated by using a haze 25 meter. The results are shown in Table 1.

EXAMPLE 2

Example 1 was repeated except that a maleic anhydride-grafted ethylene/propylene random copolymer (MAH-EPR-2; MFR₂ 1.9 g/10 min., the amount of maleic 30 anhydride grafted 0.1 g/100 g polymer, ethylene content 80 mole%, density 0.864 g/cm³, crystallinity 4%) was used instead of MAH-EPR-1. The results are shown in Table 1.

EXAMPLE 3

35 Example 1 was repeated except that a composition (MFR₂ 1.1 g/10 min., the amount of maleic

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anhydride grafted 0.25 g/100 g of polymer, ethylene content 80 mole%, density 0.865 g/10 min., crystallinity 4%) formed by mixing 50 parts by weight of MAH-EPR-1 and 50 parts by weight of an unmodified ethylene/propylene random copolymer (EPR-1; MFR₂ 2.0 g/10 min., min., ethylene content 80 mole%, density 0.864 g/cm³, crystallinity 4%) was used instead of MAH-EPR-1. The results are shown in Table 1.

EXAMPLE 4

10 Example 1 was repeated except that a composition (MFR₂ 1.4 g/10 min., the amount of maleic anhydride grafted 0.256 g/100 g of polymer, density 0.874 g/cm³, crystallinity 9%) obtained by mixing 50 parts by weight of MAH-EPR-1 and 50 parts by weight of unmodified 15 ethylene/1-butene random copolymer (EBR-1; MFR₂ 4.0 g/10 min., ethylene content 90 mole%, density 0.885 g/cm³, crystallinity 16%) was used instead of MAH-EPR-1. The results are shown in Table 1.

EXAMPLE 5

20 Example 1 was repeated except that a composition (MFR₂ 1.3 g/10 min., the amount of maleic anhydride grafted 0.25 g/100 g of polymer, density 0.874 g/cm³, crystallinity 9%) obtained by mixing 50 parts by weight of maleic anhydride-grafted ethylene/1-butene random 25 copolymer (MAH-EBR-1; MFR₂ 3.2 g/10 min., the amount of maleic anhydride grafted 0.5 g/100 g of polymer, ethylene content 90 mole%, density 0.885 g/cm³, crystallinity 14%) with 50 parts by weight of EPR-1 used in Example 3 was used instead of MAH-EPR-1. The results are shown in 30 Table 1.

EXAMPLE 6

Example 1 was repeated except that nylon 6 (trademark Aramine CM1021XF, a product of Toray Co., Ltd.) was used instead of EVOH, and the temperature of 35 the extruder for nylon was adjusted to 250°C. The results are shown in Table 1.

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EXAMPLE 7

Example 1 was repeated except that PET-G (amorphous polyethylene terephthalate-type copolymer resulting from substitution of 1,4-cyclohexanedimethanol for part 5 of the glycol component; trademark KODAR PETG6763, a product of Eastmann Chemical) was used instead of PET and a non-stretched three-layer T-die sheet was formed. The interfacial adhesion strengths and transparency of the non-stretched sheet were evaluated as in Example 1. The 10 results are shown in Table 1.

COMPARATIVE EXAMPLE 1

Example 1 was repeated except that EPR-1 used in Example 3 was used instead of MAH-EPR-1. The results are shown in Table 1.

15 COMPARATIVE EXAMPLE 2

Example 1 was repeated except that maleic anhydride-grafted LDPE (MAH-LDPE-1, MFR₂=3.0 g/10 min., the amount of maleic anhydride grafted 0.3 g/100 g of polymer, density 0.920 g/cm³, crystallinity 50%) was 20 used instead of MAH-EPR-1. The results are shown in Table 1.

COMPARATIVE EXAMPLE 3

Example 1 was repeated except that a molten mixture (MFR₂ 1.5 g/10 min., the amount of maleic 25 anhydride grafted 0.21 g/100 g of polymer, density 0.907 g/cm³, crystallinity 40%) obtained by mixing 70 parts by weight of maleic anhydride-grafted polyethylene (MAH-PE; MFR₂ 2.0 g/10 min., the amount of maleic anhydride grafted 0.3 g/100 g of polymer, density 0.925 g/cm³, 30 crystallinity 55%) and 30 parts by weight of EPR-1 used in Example 3 was used instead of MAH-EPR-1. The results are shown in Table 1.

COMPARATIVE EXAMPLE 4

Comparative Example 3 was repeated except that 35 MAH-EPR-1 used in Example 1 was used instead of EPR-1 used in Comparative Example 3. The results are also shown in Table 1.

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Table I

	Unstretched Film			Stretched Film		
	F ₁ (g/15mm width)	F ₂ (g/15mm width)	Haze (%)	F ₃ (g/15mm width)	F ₄ (g/15mm width)	Haze (%)
Example 1						
2	1200	Unpeelable	1.0	370	Unpeelable	0.6
3	1050	1270	0.9	290	380	0.5
4	1140	Unpeelable	1.0	340	450	0.6
5	970	Unpeelable	1.1	270	340	0.7
6	940	Unpeelable	1.1	320	330	0.7
7	1250	Unpeelable	0.9	380	Unpeelable	0.5
Comparative Example 1						
2	1300	Unpeelable	0.8	-	-	-
3	20	30	0.9	10	10	0.5
4	150	700	2.5	20	30	1.4
5	470	1500	4.2	40	60	2.0
6	520	Unpeelable	4.6	50	70	2.3

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CLAIMS

1. A laminated structure which comprises, in order;
 - (A) a layer of a polyester,
 - (B) a layer of a modified ethylene/alpha-olefin random copolymer, the modified copolymer containing 0.01 to 5 10% by weight, based on the trunk ethylene/alpha-olefin random copolymer, of an unsaturated carboxylic acid or derivative thereof grafted thereto, and having an ethylene content of 75 to 95 mole%, a melt flow rate of 0.1 to 50 g/10 min., a density of 0.850 to 0.905 g/cm³ and a 10 crystallinity of less than 30%, and
 - (C) a layer of a saponified olefin/vinyl acetate copolymer or a polyamide resin layer.
2. A laminated structure according to claim 1 wherein the amount of grafted unsaturated carboxylic acid or 15 derivative is 0.1 to 5% by weight.
3. A laminated structure according to claim 1 or 2 wherein the modified ethylene/alpha-olefin random copolymer has an ethylene content of 77 to 93 mole%.
4. A laminated structure according to claim 1, 2 or 3 20 wherein the modified ethylene/alpha-olefin random copolymer has a melt flow rate of 0.2 to 20 g/10 min.
5. A laminated structure according to any one of the preceding claims wherein the modified ethylene/alpha-olefin random copolymer has a density of 0.860 to 0.900 g/cm³.
- 25 6. A laminated structure according to any one of the preceding claims wherein the modified ethylene/alpha-olefin random copolymer has a crystallinity of less than 10%.
7. A laminated structure according to any one of the preceding claims wherein the polyester has units derived 30 from an aromatic dicarboxylic acid and an aliphatic alcohol.
8. A laminated structure according to any one of the preceding claims wherein the saponified olefin/vinyl acetate copolymer has an olefin content of 15 to 60 mole% and a degree of saponification of at least 50%.

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9. A laminated structure according to any one of the claims 1 to 7 wherein the polyamide resin is nylon 6.
10. A laminated structure according to any one of the preceding claims wherein the layer (A) has a thickness of 0.02 to 5 mm, the layer (B) has a thickness of 0.01 to 1 mm, and the layer (C) has a thickness of 0.01 to 1 mm.